
(12) **UK Patent Application** (19) **GB** (11) **2 065 091 A**

(21) Application No **8036097**
(22) Date of filing **10 Nov 1980**

(30) Priority data

(31) **79/39185**
80/19460

(32) **13 Nov 1979**
13 Jun 1980

(33) **United Kingdom (GB)**

(43) Application published
24 Jun 1981

(51) **INT CL³**
C01B 31/00 B01D 53/34
// A24D 3/14

(52) Domestic classification
C1A J241 J285 J290
J293 J370 J371 J383 J3
J453 J455 J462 J580
S202 S21X S21Y S432
S46X S46Y S471 S48X
S491 S492 S493 S601
S60X S60Y S610 S680
S711 SB
A2C 1E2 1E3

(56) Documents cited
GB 1235880

(58) Field of search
A2C
B1C
C1A

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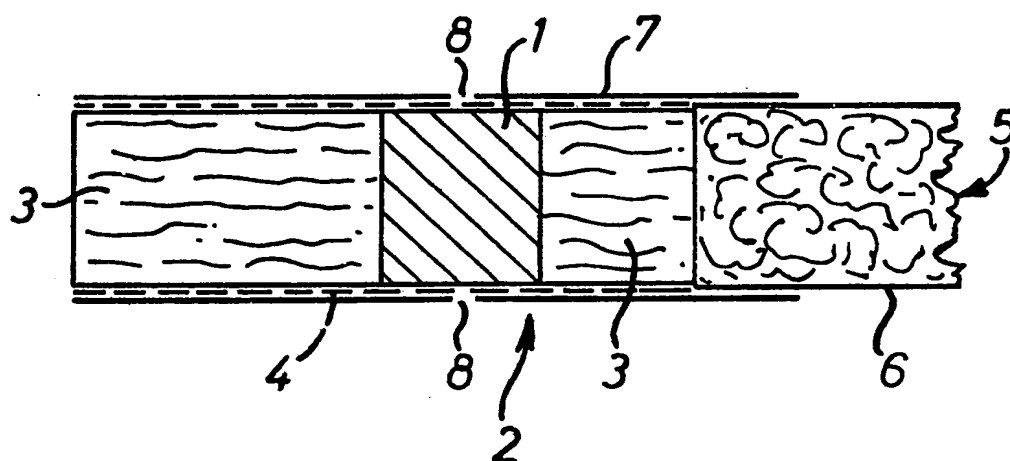
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(54) Improvements relating to
tobacco-smoke filters

(57) A nitric oxide filtration material or filter, particularly for tobacco smoke, comprises activated carbon, preferably in particulate form, upon which has been absorbed a C-nitroso compound, any substituent group of said compound, other than a nitroso group, being such as to have no critical adverse effect on the integrity of the nitroso group. The compound is preferably an aromatic compound such as nitrosobenzene, 2-nitrosotoluene or 2,4, 6-trimethylnitrosobenzene. Suitably

the carbon is coal-based. Advantageously the carbon has been loaded with a metal, such as copper or iron at a loading level in the range of 0.1% to 5% by weight on an untreated carbon basis. The loading level of the C-nitroso compound may be in the range of 2% to 15%, preferably 2% to 7% by weight also on an untreated carbon basis. Such a filter comprising the C-nitroso compound may advantageously also comprise ventilation means. The invention also comprises a method for the filtration of nitric oxide from a gaseous medium which comprises passing the medium in contact with a filtration material as hereinbefore defined.

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SPECIFICATION

Improvements relating to tobacco-smoke filters

This invention concerns improvements relating to filters or filtration materials.

It is an object of the invention to provide an improved filter or filtration material for the removal of nitric oxide from a gaseous medium, air for example.

It is a particular object of the invention to provide a filter of filtration material which is effective in practical manner for the removal of nitric oxide in smoke from cigarettes or other smoking articles.

According to the present invention, a filter of filtration material for nitric oxide removal comprises activated carbon, upon which has been adsorbed a C-nitroso compound, any substituent group or groups of said compound, other than the nitroso group or groups, being of such identity and position in the molecular structure of said compound as to have no adverse effect or no critical adverse effect on the integrity of the nitroso group or groups.

Advantageously, the C-nitroso compound is an aromatic compound.

The activated carbon may be in a fibrous, filamentary or cloth form, but is preferably in particulate form and is preferably granular coal-based carbon. The carbon should be of a porous type, so that it has a relatively high total surface area. Particle size may suitably be in a range of 5 by 50 British Standard Sieve Gauge preferably in a range of 12 by 30. The carbon may be in a bonded condition, at least in a loosely bonded condition permitting access of the smoke to the carbon.

The loading level of the C-nitroso compound on the carbon is suitably in the range 2% to 15% by weight on an untreated carbon basis and advantageously from 5% to 7%. Whereas a filter effective to remove at least 30% to 35% by weight of nitric oxide from tobacco smoke would generally be regarded as useful, with a cigarette of high nitric oxide delivery a lower filtration efficiency for nitric oxide, say as low as about 20%, could still be regarded as having utility.

The use in a filter of a C-nitroso compound adsorbed upon activated carbon may be applied in combination with filter ventilation, a synergistic effect being apparently obtainable. In some cases, accordingly, a required overall filter efficiency for nitric oxide (NO filtration efficiency) may be attainable in a practically convenient and/or economic fashion by employing the aforesaid combination.

The filtration efficiency for nitric oxide may also be significantly enhanced by employing for the activated carbon one which has been loaded with a suitable metal. For best results the metal must be in the metallic, i.e. zero valency, form. The metal may be copper or iron at a loading level on an untreated carbon basis of up to about 10% by weight. The loading level of the metal may advantageously be in a range of about 1% to 5%. Not only can NO filtration efficiencies within the range obtainable with unmetallised carbon be obtained at much reduced loading levels of C-nitroso compound by metallising the carbon, but even with modest loading levels of metal, NO filtration values can be achieved which significantly exceed those which may be feasibly achieved using unmetallised carbon. Thus with metallised carbons the loading level of the C-nitroso compound may be within a range of about 2% to about 5% on an untreated carbon basis.

Metallisation of the carbon also has the advantage that the filtration material is thereby rendered effective for the removal of a tobacco smoke component additional to nitric oxide, for instance hydrogen cyanide.

EXAMPLE I

Nitrosobenzene was prepared by the method of Coleman et al (Organic Syntheses Collective, Vol. III, edited by E. C. Horning, PP. 668-70) and a quantity of granular activated carbon, supplied under the trade name Anthrasorb CC 1236 by Cardian Chemical Co., of Cheltenham, England, thoroughly wetted with a solution in ethanol of the nitrosobenzene. The solvent was then allowed to evaporate at room temperature. The resultant loading level of the nitrosobenzene adsorbed on the carbon was 7%.

100 mg of the thus treated granular carbon was disposed in the cavity of a tripple filter comprising at each end of the cavity a 5 mm long filter plug of cellulose acetate. The filter was attached to a cigarette rod having a filter of flue-cured tobacco. The thus formed filter cigarette was then smoked under standard smoking conditions, i.e. a 35 cc volume puff of two seconds duration every minute, to a cigarette rod but length of 8 mm. It was observed that the filter removed 63% by weight of the nitric oxide from the tobacco smoke.

EXAMPLE II

The procedure of Example I was repeated using, with same loading level, 2-nitrosotoluene, obtained from Aldrich Chemical Co. Ltd., of Gillingham, Dorset, England, instead of the nitrosobenzene. The NO filtration efficiency was observed to be 50%.

EXAMPLE III

2, 4, 6-trimethylnitrosobenzene was prepared by the method of Di Nunno (Journal of Chemical Society (Section C), 1970, p. 1423). Using this so prepared compound, the procedure of Example I was again repeated. The observed NO filtration efficiency was 40%.

EXAMPLE IV

Example I was repeated using 2,4, 6-tri-*t*-butylnitrosobenzene prepared from 2, 4, 6-tri-*t*-butylaniline according to the method of Di Nunno (*ibid*), the intermediate being prepared by the method of Bartlett (Journal of American Chemical Society, 1956, Vol. 76, p. 2349). The observed NO filtration efficiency was 65%.

EXAMPLE V

4-chloronitrosobenzene was obtained from 4-chloroaniline according to the method of Di Nunno (*ibid*), and Example I was repeated. The observed NO filtration efficiency was 62%.

EXAMPLE VI

Example I was repeated except that the C-nitroso compound used was 2, 4, 6-trichloronitrosobenzene. The 2, 4, 6-trichloronitrosobenzene was obtained by the method of Di Nunno (*ibid*), the crude product being twice recrystallised from glacial acetic acid. The observed NO filtration efficiency was 38%.

EXAMPLE VII

Example I was repeated using 2-methyl-2-nitrosopropane obtained from Aldrich Chemical Co Ltd. The observed NO filtration efficiency was 33%.

EXAMPLE VIII

Example I was repeated using 2-methyl-2-nitrosopentan-2-one prepared by the method of Harries *et al* (Berichte, Vol. 3, 1898, pp. 1399 and 1808). The observed NO filtration efficiency was 59%.

EXAMPLE IX

2, 6-dimethyl-6-nitrosohept-2-en-4-one was prepared by treating 4-oxo-tetramethyl-piperidino-*l*-oxyl bromide with a mixture of aqueous sodium hydroxide and carbon tetrachloride. This C-nitroso compound was used in once again repeating Example I. The observed filtration efficiency for nitric oxide was 55%.

EXAMPLE X

Example I was repeated using; N, N-dimethyl-4-nitrosoaniline, obtained from Aldrich Chemical Co. Ltd. The observed NO filtration efficiency was 20%.

The observed NO filtration efficiency obtained using the material of Example X is low. This is because the N,N-dimethyl-amino group present in the molecular structure of the C-nitroso compound has an adverse effect on the integrity of the nitroso group. However, as mentioned above, in some cases, an NO filtration efficiency of the material as low as about 20% could be acceptable. The material could, moreover, be used in conjunction with filter ventilation. One mode of providing for such ventilation will be referred to below in connection with Example XIII.

EXAMPLE XI

Nitrosobenzene was adsorbed at various loadings on a number of different granular activated carbons and the respective nitric oxide filtration efficiencies were determined as per Example I but with cigarette rods having blended tobacco fillers. The results are set out in the Table 1 below:

Carbon \ Loading %	0	2	5	7	10	15
	Nitric Oxide Filtration Efficiency (%)					
B.P.L.	5	33	47	48	50	55
Anthrasorb CC 1236	0	25	38	39	41	47
MF3	0	11	29	29	32	38
Actibon X	2	23	34	33	38	37
Picatif 60143	8	20	30	36	37	36
207C	7	23	33	39	44	43

B. P. L. and Anthrasorb CC 1236 are anthracite carbons, the former being supplied by Pittsburgh Activated Carbon Co. of Pennsylvania, U.S.A. MF 3, Picatif 60413 and 207C are coconut based carbons supplied respectively by Chemviron Ltd., of Brussels, Belgium; Société Pica, of Paris, France; and Sutcliffe-Speakman Ltd., of Leigh, Lancashire, England. Actibon X is a wood based carbon supplied by Hooker-Mexicana S.A., of Mexico.

From the above Table it may be seen that as the loading of nitrosobenzene is increased from 0% to 5% the NO filtration efficiencies, for all of the carbons, increase rapidly, whereas when the loading levels are further increased up to 15% the NO filtration efficiencies rise more gradually. These results indicate that an optimum loading lies within an approximate range of 5% - 7%.

10 EXAMPLE XII

Example 1 was repeated using nitrosobenzene at a loading level of 5% on Anthrasorb CC 1844 granular activated carbon. The cigarette rod has a filler of blended tobacco. The observed NO filtration efficiency was 53%.

15 EXAMPLE XIII

Nitrosobenzene was adsorbed at a 5% loading level on B.P.L. granular activated carbon. a 100 mg amount of the treated carbon was disposed in the cavity 1 of each of a number of triple filters 2 having at each end of the cavity 1, a cellulose acetate filter plug 3, as illustrated by the single figure of the accompanying drawing, which is a diagrammatic longitudinal section through a filter. Each filter, enclosed in a porous plugwrap 4, was attached to a cigarette rod 5 having a filler of blended tobacco within a wrapper 6. The filters 2, otherwise identical, were attached to respective rods 5 by tippings 7 provided with varying numbers of rows of laser-formed microperforations 8 disposed around a central region of the cavity 1 to give varying degrees of ventilation.

The thus formed filter cigarettes, and control cigarettes, of similar delivery, were smoked under standard smoking conditions to a cigarette-rod butt length of 8 mm and the nitric oxide delivery for each cigarette was determined. The control cigarettes varied from the above-described cigarettes in that the cavities of the filters thereof contained 100 mg of untreated B.P.L. granular activated carbon.

The results are set out in Table 2 below:

Ventilation (%)	Untreated BPL Carbon		BPL Impregnated with 5% Nitrosobenzene		Retention due to Nitrosobenzene (%)	Expected Retention due to Nitrosobenzene (%)
	Delivery ($\mu\text{g}/\text{cig}$)	Reduction due to Ventilation (%)	Delivery ($\mu\text{g}/\text{cig}$)	Total Reduction (%) due to vent ⁿ + Nitrosobenzene		
0	210	—	123	41	41	42
23	148	30	74	65	50	46
43	118	44	41	80	65	48
58	88	58	21	90	76	50
59	77	63	19	91	75	51

The values for expected nitric oxide retention due to the nitrosobenzene were derived from a relationship between retention and nitric oxide delivery determined by attaching non-ventilated triple filters, each containing 100 mg of B.P.L. granular activated carbon carrying a 5% loading of nitrosobenzene, to cigarette rods of different, predetermined, nitric oxide deliveries. The relationship was found to be a virtually straight-line relationship.

A comparison of the actual and expected values of nitric oxide retention shows that with increasing ventilation the actual retention is increasingly higher than the expected retention. That is to say, a synergistic effect is obtained.

A similar synergistic effect on nitric oxide retention was observed when the ventilation perforations were positioned over the mouth end cellulose acetate plugs instead of over the cavities. When the perforations were positioned over the tobacco end plugs the synergism was less pronounced.

EXAMPLE XIV

B.P.L. granular activated carbons with zero, 0.1%, 1%, 5% and 10% loading levels of copper were impregnated with nitrosobenzene at loading levels of zero, 1%, 2%, 5%, 10% and 15% on a metallised carbon basis.

- 5 100 mg amounts of the respective carbons were disposed in the cavities of triple filters, each filter comprising at each end of the cavity a 5 mm long plug of cellulose acetate. The filters were attached to cigarette rods having a filler of blended tobacco. Each cigarette rod was such as to give a nitric oxide delivery of about 200 μ g when no filter was attached. The thus formed filter cigarettes were then smoked under standard smoking conditions. In each case a determination was made of the NO filtration efficiency of the filter. The results obtained are shown in Table 3 below:

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	Loading Copper %				
	0	0.1	1.0	5.0	10.0
Loading Nitrosobenzene (%)	Nitric Oxide Filtration Efficiencies (%)				
0	<10	<10	<10	<10	<10
1.0	17	25	37	33	26
2.0	36	40	48	48	41
5.0	42	46	62.5	65	54.5
10.0	43	56.5	65	61	54
15.0	47	58	67.5	67	53

- From Table 3 it may be seen that even a loading level of copper as low as 0.1% improves the NO filtration efficiency at all of the loading levels of nitrosobenzene. A 1.0% loading of copper results in even more significantly enhanced NO filtration efficiencies. An increase in copper loading to a 5% level results in NO filtration efficiencies closely similar to those obtained at a 1% loading. An increase to a 10% loading results in NO filtration efficiencies lower than those obtained at the 1% and 5% loadings and, for higher nitrosobenzene loadings, the filtration efficiencies are even lower than those obtained with 0.1% copper. It may thus be deduced that when using nitrosobenzene as the C-nitroso compound and selecting B.P.L. as the activated carbon, optimum copper loading levels are in a range of about 1% to about 5%. It may also be observed that at any particular loading level of copper there is a steep rise in the NO filtration efficiency on increasing the loading level of nitrosobenzene from zero to 2%, but that only relatively small increases in NO filtration efficiency result from going from 5% to 10% and from 10% to 15%, in nitrosobenzene loading levels. Hence there is no apparent benefit in using loading levels of nitrosobenzene greater than about 5% particularly when the copper loading level is in the range of about 1% to 5%.

EXAMPLE XV

The procedure of Example XIV was followed except that activated carbons loaded with iron were

used, the carbons again being of B.P.L. type. The results obtained with the iron-loaded carbons are shown in Table 4 below:

	Loading Iron %				
	0	0.1	1.0	5.0	10.0
Loading Nitrosobenzene (%)	Nitric Oxide Filtration Efficiencies (%)				
0	<10	<10	<10	<10	<10
1.0	17	21	26	25	25
2.0	36	30	41	37	40
5.0	42	40	50	48	44
10.0	43	45	58	50	42
15.0	47	52	61	54	51

As may be seen Table 4 the general pattern of results is similar to that obtained using the copper-loaded carbons, although the enhancement in NO filtration efficiencies over non-metallised carbon is not quite so pronounced as was the case with copper metallisation. The results do though illustrate an effect of very definite utility especially when the relatively lower cost of iron-loaded carbons is taken into account. It may also be observed from Table 4 that the NO filtration efficiencies at a 5% loading level of iron are not closely similar to the filtration efficiencies at a 1% loading level, as was the case with copper loaded carbons, but are somewhat lower. Thus it may be concluded that with nitrosobenzene-treated B.P.L. carbons optimum loadings of iron are in the region of the 1% level.

Although in the above examples the C-nitroso treated carbons are deposited in the cavities of triple filters, they may, of course, be otherwise incorporated in cigarette filters. For example, a C-nitroso treated carbon could be dispersed throughout a plug of cellulose acetate or other fibrous filtration material.

Provision for filter ventilation, for example as discussed with reference to Example XIII, may be utilised also in conjunction with filters in accordance with any of the above Examples.

CLAIMS

1. A nitric oxide filtration material comprising activated carbon upon which has been adsorbed a C-nitroso compound, any substituent group or groups of said compound, other than the nitroso group or groups, being of such identity and position in the molecular structure of said compound as to have no adverse effect or no critical adverse effect on the integrity of the nitroso group or groups.

2. A filtration material according to Claim 1, wherein the C-nitroso compound is an aromatic compound.

3. A filtration material according to Claim 2, wherein the C-nitroso compound is nitrosobenzene.

4. A filtration material according to Claim 2, wherein the C-nitroso compound is 2-nitrosotoluene.

5. A filtration material according to Claim 2, wherein the C-nitroso compound is 2, 4, 6-trimethyl-nitrosobenzene.

6. A filtration material according to any preceding claim, wherein the activated carbon is a coal-based carbon.

7. A filtration material according to any preceding claim, wherein the activated carbon is in particulate form.

8. A filtration material according to any preceding claim, wherein the activated carbon has been loaded with a metal.

9. A filtration material according to Claim 8, wherein the metal is copper or iron.

10. A filtration material according to Claim 8 or 9, wherein the loading level of the metal is in the range of 0.1% to 5% by weight on an untreated carbon basis.

11. A filtration material according to any preceding claim, wherein the loading level of the C-nitroso compound is in the range of 2% to 15% by weight on an untreated carbon basis.

12. A filtration material according to Claim 11, wherein the loading level of the C-nitroso compound is in the range of 2% to 7%.

13. A filtration material substantially as hereinbefore described with reference to any one of the Examples.

14. A filter comprising a filtration material according to any preceding claim.

15. A smoking article filter comprising a filtration material according to any one of claims 1 to 13.

5 16. A filter according to Claim 15, said material being disposed in the filter and the filter comprising ventilation means arranged to provide air into the filter. 5

17. A method for the filtration of nitric oxide from a gaseous medium which comprises passing the said medium in contact with a filtration material according to any one of Claims 1 to 13.

Printed for Her Majesty's Stationery Office by the Courier Press, Leamington Spa, 1981. Published by the Patent Office,
25 Southampton Buildings, London, WC2A 1AY, from which copies may be obtained.